



CRF INC. EXPANDS CLINICAL ADVISORY BOARD

- Respected clinical and scientific advisors provide scientific and therapeutic area expertise for customer clinical programs -

Waltham, MA – June 18, 2008: CRF Inc., the leading global provider of electronic Patient Reported Outcomes (ePRO) and wireless data collection solutions for the Life Science Industry, today announced the expansion of its world-class Clinical Advisory Board. Comprised of leading clinical and scientific experts, the Clinical Advisory Board will play an important role in providing specialist scientific and therapeutic advice for customers' clinical programs. In addition, the board will provide advice and guidance on the interpretation of the FDA's Draft Guidance for the Industry on Patient Reported Outcomes and validation requirements.

"We are privileged to bring together experts from some of the leading clinical and research institutes around the world," said Pamela McNamara, Chief Executive Officer at CRF Inc. "Each member of our Clinical Advisory Board brings a wealth of knowledge and experience that will be instrumental in assisting our customers with their clinical programs. We are thrilled to welcome these clinical and scientific experts to CRF."

CRF's Clinical Advisory Board is comprised of renowned thought leaders in the fields of instrument development; psychometric validation; clinical study design and execution; and patient behavior as well as expertise in specific therapeutic areas. Rachael King, CRF Inc. Vice President for European Operations chairs the Clinical Advisory Board. Ms. King has an MA in Pharmacology from Cambridge University (UK) and 12 years experience working for a major Pharmaceutical company plus 14 years in the ePRO space. The Clinical Advisory Board includes Nancy Kline Leidy, PhD from the United BioSource Corporation; Richard Lennox, PhD of Psychometric Technologies; Ville Ojanen, PhD of Cogiteq Ltd.; and Susan MacIntyre RN, BSN of US teaching and Research Institutions, NHS Research Centres specializing in Sleep and Neurology.



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About CRF Inc.

CRF Inc. (CRF) is a leading global provider of electronic Patient Reported Outcomes (ePRO) and wireless data collection solutions for the Life Science Industry. Through innovative technology and a thorough understanding of drug development and mobile computing, the company is driving the change to safer and more efficient, paper-free clinical trials. CRF's technology has been used by more than 180,000 patients across 59 countries in 60 languages for 45 indications. The company has demonstrated one of the industry's highest patient compliance rates - an average 95% compliance through Phase I, II, III and IV clinical trials.

CRF's award-winning product, the TrialMax[®] suite, is a flexible and configurable ePRO technology that provides real-time patient monitoring, outstanding data accuracy and increased safety through compliance. The TrialMax[®] application's unique features enable clinical trial sponsors to rapidly collect valuable data and conduct complex clinical trials with greater flexibility than other ePRO solutions. CRF's experience, combined with its dedication to achieving high quality and responsive customer service, has made the company one of the life science industry's most trusted partners. For more information, please visit www.crfhealth.com.

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